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38-CV-2021-900350.00

CIRCUIT COURT OF

HOUSTON COUNTY, ALABAMA

CARLA H. WOODALL, CLERK

#### IN THE CIRCUIT COURT OF HOUSTON COUNTY, ALABAMA

WALTER JOHNSON,	)	
Plaintiff,	)	
v.	)	
3M COMPANY, INC.; and FICTITIOUS DEFENDANTS "A-Z," are those other persons, firms, and/or corporations whose negligence or wantonness caused and/or contributed to cause the injuries sustained by Plaintiff, whose identity has not yet been determined but whose names will be	) ) Case No ) ) ) )	
substituted when ascertained,		
Defendants.	)	

#### **COMPLAINT**

#### **Statement of the Parties**

- 1. Plaintiff Walter Johnson, hereinafter referred to as "Johnson" and/or "Plaintiff", is over the age of nineteen and is a resident of Houston County, Alabama.
- 2. Defendant 3M Company, Inc., hereinafter referred to as "3M" and/or "Defendant", is a foreign corporation doing business in the state of Alabama.
- 3. Fictitious Defendants "A–Z," whether singular or plural, are those other persons, firms, and/or corporations whose negligence or wantonness caused and/or contributed to cause the injuries sustained by Plaintiff, whose identity has not yet been determined but whose names will be substituted when ascertained.
- 4. When reference is made in this Complaint to any act or omission of any of the Defendants, it shall be deemed that the officers, directors, agents, employees, or representatives of the Defendants committed or authorized such act or omission, or failed to adequately supervise or properly control or direct their employees while engaged in the management,

direction, operation, or control of the affairs of Defendants, and did so while acting within the scope of their duties, employment or agency.

5. The term "Defendants" is made to refer to all real and Fictitious Defendants described in the style of this Complaint.

#### **Statement of Facts**

- 6. Plaintiff brings this action for damages for personal injury resulting from exposure to aqueous film-forming foams ("AFFF") containing the toxic chemicals collectively known as per and polyfluoroalkyl substances ("PFAS"). PFAS includes, but is not limited to, perfluorooctanoic acid ("PFOA") and perfluorooctane sulfonic acid ("PFOS") and related chemicals including those that degrade to PFOA and/or PFOS. AFFF is a Class-B firefighting foam. It is mixed with water and used to extinguish fires that are difficult to fight, particularly those that involve petroleum or other flammable liquids.
- 7. Plaintiff began his employment as a firefighter with the Dothan Fire Department in 1997 and was first exposed to AFFF during training that same year.
- 8. Plaintiff was continuously exposed to AFFF throughout his career until his retirement in December 2019.
  - 9. Plaintiff was diagnosed with multiple myeloma in September 2019.
- Defendants designed, marketed, developed, manufactured, distributed, released, trained users, produced instructional materials, promoted, sold, and/or otherwise released into the stream of commerce AFFF with knowledge that it contained highly toxic and bio persistent PFAS, which would expose end users of the product to the risks associated with PFAS.
- Defendants designed, marketed, developed, manufactured, distributed, released, trained users on, produced instructional materials for, sold, and/or otherwise handled and/or used AFFF containing PFAS, in such a way as to cause the contamination of Plaintiff's blood and/or

body with PFAS, and the resultant biopersistence and bioaccumulation of such PFAS in the blood and/or body of Plaintiff.

- 12. As a result of Defendants' conduct, Plaintiff was injured as follows:
  - (a) He was diagnosed with multiple myeloma;
  - (b) He suffered physical pain and will continue to do so in the future;
  - (c) He suffered mental anguish and will continue to do so in the future;
  - (d) He is permanently injured and damaged;
  - (e) He has incurred medical expenses and will do so in the future; and
  - (f) He has been otherwise injured and damaged.

## COUNT ONE (Negligence)

- 13. Plaintiff hereby incorporates by reference the allegations contained in the preceding paragraphs of this Complaint as if restated in full herein.
- 14. Defendants had a duty to individuals, including the Plaintiff, to exercise reasonable ordinary, and appropriate care in the manufacturing, design, labeling, packaging, testing, instruction, warning, selling, marketing, distribution, and training related to the AFFF product.
- 15. Defendants breached their duty of care and were negligent, grossly negligent, reckless and willful as described herein in the design, manufacture, labeling, warning, instruction, training, selling, marketing, and distribution of the AFFF products in one or more of the following respects:
  - a. Failing to design the products so as to avoid an unreasonable risk of harm to individuals, including the Plaintiff;
  - b. Failing to use reasonable care in the testing of the products so as to avoid an unreasonable risk of harm to individuals, including the Plaintiff;

- c. Failing to use appropriate care in inspecting the products so as to avoid an unreasonable risk of harm to individuals, including the Plaintiff;
- d. Failing to use appropriate care in instructing and/or warning the public as set forth herein of risks associated with the products, so as to avoid unreasonable risk of harm to individuals, including the Plaintiff;
- e. Failing to use reasonable care in marketing, promoting, and advertising the products so as to avoid unreasonable risk of harm to individuals, including the Plaintiff;
- f. Otherwise negligently or carelessly designing, manufacturing, marketing, distributing, warning; and
- g. In selling and or distributing a product which was inherently dangerous to the public.
- 16. As a direct and proximate result of Defendants' negligence, the Plaintiff has been injured, sustained severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort, economic loss and damages including, but not limited to medical expenses, lost income, and/or other damages.

### COUNT TWO (Battery)

- 17. Plaintiff hereby incorporates by reference the allegations contained in the preceding paragraphs of this Complaint as if restated in full herein.
- 18. At all relevant times, Defendants possessed knowledge that the AFFF containing PFAS which they designed, engineered, manufactured, fabricated, sold, handled, released, trained users on, produced instructional materials for, used, and/or distributed were biopersistent,

bio-accumulative, toxic, potentially carcinogenic, and/or otherwise harmful/injurious and that

their continued manufacture, use, sale, handling, release, and distribution would result in Plaintiff having PFAS in Plaintiff's blood, and the biopersistence and bioaccumulation of such PFAS in Plaintiff's blood.

- 19. However, despite possessing such knowledge, Defendants knowingly, purposefully, and/or intentionally continued to engage in such acts and/or omissions, including but not limited to all such acts and/or omissions described in this Complaint, that continued to result in Plaintiff accumulating PFAS in Plaintiff's blood and/or body, and such PFAS persisting and accumulating in Plaintiff's blood and/or body.
- 20. Defendants did not seek or obtain permission or consent from Plaintiff to put or allow PFAS materials into Plaintiff's blood and/or body, or to persist in and/or accumulate in Plaintiff's blood and/or body.
- 21. Entry into, persistence in, and accumulation of such PFAS in Plaintiff's body and/or blood without permission or consent is an unlawful and harmful and/or offensive physical invasion and/or contact with Plaintiff's person and unreasonably interferes with Plaintiff's rightful use and possession of Plaintiff's blood and/or body.
- 22. At all relevant times, the PFAS present in the blood of Plaintiff originated from Defendants' acts and/or omissions.
- 23. Defendants continue to knowingly, intentionally, and/or purposefully engage in acts and/or omissions that result in the unlawful and unconsented-to physical invasion and/or contact with Plaintiff that resulted in persisting and accumulating levels of PFAS in Plaintiff's blood.
- 24. Plaintiff, and any reasonable person, would find the contact at issue harmful and/or offensive.

- 25. Defendants acted intentionally with the knowledge and/or belief that the contact, presence and/or invasion of PFAS with, onto and/or into Plaintiff's blood serum, including its persistence and accumulation in such serum, was substantially certain to result from those very acts and/or omissions.
- 26. Defendants' intentional acts and/or omissions resulted directly and/or indirectly in harmful contact with Plaintiff's blood and/or body.
- 27. The continued presence, persistence, and accumulation of PFAS in the blood and/or body of Plaintiff is offensive, unreasonable, and/or harmful, and thereby constitutes a battery.
- 28. The presence of PFAS in the blood and/or body of Plaintiff altered the structure and/or function of such blood and/or body parts and resulted in cancer.
- 29. As a direct and proximate result of the foregoing acts and omissions, Plaintiff suffered physical injury for which Defendants are therefore liable.

## COUNT THREE (Inadequate Warning)

- 30. Plaintiff hereby incorporates by reference the allegations contained in the preceding paragraphs of this Complaint as if restated in full herein.
  - 31. Defendants knew or should have known:
    - a. exposure to AFFF containing PFAS was hazardous to human health;
    - b. the manner in which they were designing, marketing, developing, manufacturing, distributing, releasing, training, instructing, promoting, and selling AFFF containing PFAS was hazardous to human health; and

- c. the manner in which they were designing, marketing, developing, manufacturing, marketing, distributing, releasing, training, instructing, promotion and selling AFFF containing PFAS would result in the contamination of Plaintiff's blood and/or body as a result of exposure.
- 32. Defendants had a duty to warn of the hazards associated with AFFF containing PFAS entering the blood and/or body of Plaintiff because they knew of the dangerous, hazardous, and toxic properties of AFFF containing PFAS. Defendants failed to provide sufficient warning to purchasers that the use of their AFFF products would cause PFAS to be released and cause the exposure and bioaccumulation of these toxic chemicals in the blood and/or body of Plaintiff.
- 33. Adequate instructions and warnings on the AFFF containing PFAS could have reduced or avoided these foreseeable risks of harm and injury to Plaintiff. If Defendants provided adequate warnings:
  - a. Plaintiff could have and would have taken measures to avoid or lessen exposure; and
  - b. end users and governments could have taken steps to reduce or prevent the release of PFASs into the blood and/or body of Plaintiff. Defendants' failure to warn was a direct and proximate cause of Plaintiff's injuries from PFAS that came from the use, storage, and disposal of AFFF containing PFAS. Crucially, Defendants' failure to provide adequate and sufficient warnings for the AFFF containing PFAS they designed, marketed, manufactured, distributed, released, promoted, and sold renders the AFFF a defective product.
- 34. Defendants were negligent in their failure to provide Plaintiff with adequate warnings or instruction that the use of their AFFF products would cause PFAS to be released into the blood and/or body of Plaintiff. As a result of Defendants' conduct and the resulting contamination, Plaintiff suffered severe personal injuries by exposure to AFFF containing PFAS.
- 35. Defendants' negligent failure to warn directly and proximately caused the harm to and damages suffered by Plaintiff.

### COUNT FOUR (Design Defect)

- 36. Plaintiff hereby incorporates by reference the allegations contained in the preceding paragraphs of this Complaint as if restated in full herein.
  - 37. Defendants knew or should have known:
    - a. exposure to AFFF containing PFAS is hazardous to human health;
    - b. he manner in which AFFF containing PFAS was designed, manufactured, marketed, distributed, and sold was hazardous to human health; and
    - C. the manner in which AFFF containing PFAS was designed, manufactured, marketed, distributed, and could and would release PFAS into Plaintiff and cause the exposure and bioaccumulation of these toxic and poisonous chemicals in the blood and/or body of Plaintiff.
- 38. Knowing of the dangerous and hazardous properties of the AFFF containing PFAS, Defendants could have designed, manufactured, marketed, distributed, and sold alternative designs or formulations of AFFF that did not contain hazardous and toxic PFAS. These alternative designs and formulations were already available, practical, and technologically feasible. The use of these alternative designs would have reduced or prevented reasonably foreseeable harm to Plaintiff caused by the Defendants' design, manufacture, marketing, distribution, and sale of AFFF containing hazardous and toxic PFAS.
- 39. The AFFF containing PFAS that was designed, manufactured, marketed, distributed, and sold by the Defendants was so hazardous, toxic, and dangerous to human health that the act of designing, formulating, manufacturing, marketing, distributing, and selling this AFFF was unreasonably dangerous under the circumstances.

- 40. The AFFF designed, formulated, manufactured, marketed, distributed, and sold by Defendants was defectively designed and the foreseeable risk of harm could and would have been reduced or eliminated by the adoption of a reasonable alternative design that was not unreasonably dangerous. Defendants' defective design and formulation of AFFF containing PFAS was a direct and proximate cause of the contamination of the blood and/or body of Plaintiff and the persistence and accumulation of PFAS in Plaintiff's blood and/or body.
- 41. Defendants' defective design and formulation of AFFF containing PFAS caused the contamination described herein resulting in personal injuries to Plaintiff. As a direct result of the harm and injury caused by Defendants' defective design and the contamination described herein, Plaintiff has been exposed to AFFF containing PFAS and other toxic substances and has developed cancer.
- 42. Defendants' negligent failure to design a reasonably safe product directly and proximately caused the harm to and damages suffered by Plaintiff.

### **COUNT FIVE** (Strict Liability – Statutory)

- 43. Plaintiff hereby incorporates by reference the allegations contained in the preceding paragraphs of this Complaint as if restated in full herein.
- 44. Plaintiff asserts any and all remedies available under statutory causes of action from Plaintiff's state for strict liability against each Defendant.
- 45. The Defendants were engaged in designing, manufacturing, marketing, selling, and distribution of AFFF.

- 46. AFFF was in a defective condition and unreasonably dangerous to users and/or consumers when designed, manufactured, marketed, sold, and/or distributed to the public by the Defendants.
- 47. As a direct and proximate result of the Defendants products' aforementioned defects, the Plaintiff has been injured, sustained severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort, economic loss and damages including, but not limited to medical expenses, lost income, and other damages.
- 48. The Defendants are strictly liable in tort to the Plaintiff for their wrongful conduct.

#### COUNT SIX (Strict Liability – Restatement)

- 49. Plaintiff hereby incorporates by reference the allegations contained in the preceding paragraphs of this Complaint as if restated in full herein.
- 50. The Plaintiff brings strict product liability claims under the common law, Section 402A of the Restatement of Torts (Second), and/or Restatement of Torts (Third) against Defendants.
- 51. As designed, manufactured, marketed, tested, assembled, equipped, distributed and/or sold by the Defendants the AFFF product was in a defective and unreasonably dangerous condition when put to reasonably anticipated use to foreseeable consumers and users, including the Plaintiff.

- 52. The Defendants had available reasonable alternative designs which would have made the AFFF product safer and would have most likely prevented the injuries and damages to the Plaintiff, thus violating state law and the Restatement of Torts.
- 53. The Defendants failed to properly and adequately warn and instruct the Plaintiff as to the proper safety and use of the Defendants product.
- 54. The Defendants failed to properly and adequately warn and instruct the Plaintiff regarding the inadequate research and testing of the product.
- 55. The Defendants' products are inherently dangerous and defective, unfit and unsafe for their intended and reasonably foreseeable uses, and do not meet or perform to the expectations.
- 56. As a proximate result of the Defendants' design, manufacture, marketing, sale, and distribution of the products, the Plaintiff has been injured and sustained severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort, and consortium, and economic damages.
- 57. By reason of the foregoing, the Defendants are strictly liable for the injuries and damages suffered by the Plaintiff, caused by these defects in the AFFF product.

### COUNT SEVEN (Fraudulent Concealment)

58. Plaintiff hereby incorporates by reference the allegations contained in the preceding paragraphs of this Complaint as if restated in full herein.

- 59. Throughout the relevant time period, Defendants knew that their products were defective and unreasonably unsafe for their intended purpose.
- 60. Defendants fraudulently concealed from and/or failed to disclose to or warn the Plaintiff, and the public that their products were defective, unsafe, and unfit for the purposes intended, and that they were not of merchantable quality.
- 61. Defendants were under a duty to the Plaintiff and the public to disclose and warn of the defective and harmful nature of the products because:
  - a. Defendants were in a superior position to know the true quality, safety and efficacy of the Defendants' products;
  - b. Defendants knowingly made false claims about the safety and quality of the Defendants' product in documents and marketing materials; and
  - c. Defendants fraudulently and affirmatively concealed the defective nature of the Defendants' products from the Plaintiff.
- 62. The facts concealed and/or not disclosed by Defendants to the Plaintiff were material facts that a reasonable person would have considered to be important in deciding whether or not to purchase and/or use the Defendants' products.
- 63. Defendants intentionally concealed and/or failed to disclose the true defective nature of the products so that the Plaintiff would use the Defendants' products, the Plaintiff justifiably acted or relied upon, to Plaintiff's detriment, the concealed and/or non-disclosed facts as evidenced by Plaintiff's use of the Defendants' products.
- 64. Defendants, by concealment or other action, intentionally prevented the Plaintiff from acquiring material information regarding the lack of safety and effectiveness of the Defendants' products and are subject to the same liability to the Plaintiff for Plaintiff's pecuniary losses, as though Defendants had stated the non-existence of such material information regarding the Defendants' products' lack of safety and effectiveness and dangers and defects, and as

though Defendants had affirmatively stated the non-existence of such matters that the Plaintiff was thus prevented from discovering the truth. Defendants therefore have liability for fraudulent concealment under all applicable laws, including, inter alia, Restatement (Second) of Torts §550 (1977).

65. As a proximate result of Defendants' conduct, the Plaintiff has been injured, and sustained severe and permanent pain, suffering, disability, impairment, loss of enjoyment of life, loss of care, comfort, and economic damages.

WHEREFORE, the Plaintiff prays judgments against the Defendants for actual, compensatory, consequential, and punitive damages, together with the costs of this action, and for such other and further relief as this Court may deem fit, just, and proper.

## COUNT EIGHT (Breach of Express and Implied Warranties)

- 66. Plaintiff hereby incorporates by reference the allegations contained in the preceding paragraphs of this Complaint as if restated in full herein.
- 67. At all times relevant hereto, the Defendants manufactured, marketed, labeled, and sold the AFFF products that has been previously alleged and described herein.
- 68. At the time the Defendants designed, developed, marketed, sold, labeled, and distributed the AFFF products, the Defendants knew of the use for which it was intended, and implied and/or expressly warranted that the product was merchantable, safe, and fit for its intended purpose.
- 69. The Defendants warranted that the product was merchantable and fit for the particular purpose for which it was intended and would be reasonably safe. These warranties were breached, and such breach proximately resulted in the injuries and damages suffered by the Plaintiff.

- 70. The Plaintiff is within the class of foreseeable users and reasonably relied upon Defendants' judgment, and the implied and/or express warranties in using the products.
- 71. The Defendants breached their implied and/or express warranties and did not meet the expectations for the performance of the product when used for its intended use and was neither of merchantable quality nor safe for its intended use in that the product has a propensity to cause serious injury, pain, and cancer.

# COUNT NINE (Wantonness)

- 72. Plaintiff hereby incorporates by reference the allegations contained in the preceding paragraphs of this Complaint as if restated in full herein.
- 73. Defendants and their employees, agents, officers, and representatives owed a duty of care to end users of their AFFF products, including Plaintiff.
  - 74. Defendants breached the duty of care owed to the Plaintiff.
- 75. The actions of Defendants and their employees, agents, officers, and representatives were willful and wanton and exhibited a reckless disregard for the life, health, and safety of the end users of Defendants' AFFF products, including Plaintiff.
- 76. As a proximate and foreseeable consequent of the actions of Defendants, Plaintiff was exposed to unreasonably dangerous toxic PFAS containing AFFF, which caused Plaintiff's injury.

RESPECTFULLY submitted this the 27<sup>th</sup> day of August, 2021.

<u>/s/ Dustin J. Fowler</u>
One of the Attorneys for Plaintiff

#### **OF COUNSEL:**

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PLAINTIFF RESPECTFULLY DEMANDS A TRIAL BY JURY

/s Dustin J. Fowler
Dustin J. Fowler

### **DEFENDANT TO BE SERVED BY CERTIFIED MAIL:**

3M Company, Inc. c/o Corporation Service Company, Inc. 641 South Lawrence Street Montgomery, AL 36104